



TITLE:

Comprehensive evaluation of airway involvement in pulmonary sarcoidosis

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Comprehensive evaluation of airway involvement in pulmonary sarcoidosis

To the Editor:

Sarcoidosis most commonly affects the thoracic lymph nodes, lung parenchyma and airway in nearly two-thirds of patients [1]. We previously reported that airflow limitation in Japanese outpatients was associated with radiographic stage IV disease, older age, smoking and bronchovascular bundle thickening on high-resolution computed tomography (HRCT) [2]. Airway hyperresponsiveness (AHR) can be another potential predisposing factor of airflow limitation [1, 3]. Impulse oscillometry (IOS) is an effort-independent and noninvasive method of assessing respiratory physiology and may detect more subtle changes than spirometry [4]. Thus, a multidisciplinary approach including AHR provocation test and IOS may provide novel insights into airway involvement of sarcoidosis compared to classical spirometric assessments. We conducted a prospective observational study to comprehensively evaluate airway involvement in patients with pulmonary sarcoidosis and investigate the impacts of various aspects of airway involvement on the long-term outcome. This study was approved by the Ethics Committee of Kyoto University (Kyoto, University; institutional board number E-530), with all subjects granting written informed consent.

We enrolled a total of 45 consecutive patients newly diagnosed with pulmonary sarcoidosis at Kyoto University Hospital between September 2008 and February 2012 for investigation. Grounds for exclusion were as follows: 1) systemic corticosteroid therapy within the preceding month; 2) history of adult asthma; 3) any other chronic lower airway disease, such as chronic obstructive pulmonary disease and bronchiectasis; 4) upper or lower airway infection within the preceding 4 weeks; 5) active neoplasm; or 6) connective tissue disease. The diagnosis of sarcoidosis was biopsy proven in 18 (40%) patients and based on a combination of clinical and laboratory findings in the remaining 27 patients [5].

At the time of diagnosis, all subjects underwent chest radiography, measurement of serum angiotensin-converting enzyme, soluble interleukin-2 receptor and Krebs von den Lungen-6 and total immunoglobulin E, pulmonary function tests (PFTs), IOS (Masterscreen IOS-J; Jaeger, Wurzburg, Germany) [4], exhaled nitric oxide fraction (F_{eNO}) measurement at expiratory flow rates of $50 \text{ mL}\cdot\text{s}^{-1}$ (NOA 280; Sievers, Boulder, CO, USA) [6] and AHR provocation test [7, 8]. HRCT of the chest (Aquilion 64; Toshiba, Tokyo, Japan) and bronchoalveolar lavage (BAL) were also performed in 44 (98%) and 36 (80%) patients, respectively [5]. The extent of organ involvement was assessed by the ACCESS (A Case-Control Epidemiologic Study of Sarcoidosis) Organ Involvement Index [9]. The AHR test was performed according to the single concentration four-step dosimeter protocol for methacholine [7, 8]. A single concentration of methacholine-chloride ($16 \text{ mg}\cdot\text{mL}^{-1}$, 1.6%) was administered through a nebuliser (MedicAid Pro Nebulizer; Jaeger) at the output of $240 \mu\text{L}\cdot\text{min}^{-1}$. In IOS, rectangular mechanical impulses in the frequency range of 5–35 Hz were applied to the respiratory system through a mouthpiece during spontaneous breathing. Resultant pressure and flow signals were analysed to determine respiratory resistance values at 5 Hz and 20 Hz (R_5 and R_{20} , respectively), reactance at 5 Hz (X_5) and area of reactance (AX) [4, 10]. HRCT images (0.5-mm slices) were analysed to determine airway wall area of the right apical bronchus and tracheal luminal area at the upper margin of aortic arch, using a modified in-house software as previously described [11].

Of the 45 patients enrolled, AHR was detected in eight (18%) out of 45 patients. Clinical manifestations, serum biomarkers, IOS, PFT, F_{eNO} , HRCT indices and BAL findings were compared between patients with



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and without AHR (table 1). Respiratory symptoms (cough, sputum production, wheezing or dyspnoea on exertion) were more prevalent in AHR-positive patients. Among IOS indices, R5 and R20 were significantly higher in AHR-positive patients, whereas R5–R20 and X5 values were similar, regardless of

TABLE 1 Comparison between patients with and without airway hyperreactivity (AHR)

AHR	Positive	Negative	p-value
Subjects	n=8	n=37	
Age years	39.5 [26.8–70.3]	51 [35.5–60.0]	NS
Males	3 [38]	16 [43]	NS
Height cm	157 [149–174]	162 [157–170]	NS
Weight kg	52.9 [45.3–78.5]	58.0 [51.3–63.0]	NS
Current or past smoker	3 [38]	20 [54]	NS
Smoking history pack-years	0 [0–10]	1.5 [0–37.4]	
Histologic confirmation	4 [50]	14 [38]	NS
Extrapulmonary involvement	5 [65]	24 [63]	NS
Radiographic stage I/II/III/IV	2 [25]/6 [75]/0 [0]/0 [0]	11 [30]/23 [62]/3 [8]/0 [0]	NS
Respiratory symptoms	6 [75]	10 [27]	0.02
mMRC[#] 1/2/3/4/5	5 [63]/2 [25]/1 [13]/0 [0]/0 [0]	31 [84]/5 [14]/1 [3]/0 [0]/0 [0]	NS
Serum biomarkers			
ACE IU·L ⁻¹	18.9 [12.5–25.9]	16.7 [14.0–26.3]	NS
sIL-2R U·mL ⁻¹	904 [710–1410]	532 [409–889]	NS
KL-6 U·mL ⁻¹	341 [251–451]	271 [214–361]	NS
IgE IU·mL ⁻¹	110 [100–370]	92 [26–338]	NS
Impulse oscillometry			
R5 kPa·L ⁻¹ ·s ⁻¹	0.33 [0.28–0.43]	0.23 [0.19–0.31]	0.01
R20 kPa·L ⁻¹ ·s ⁻¹	0.29 [0.24–0.35]	0.21 [0.17–0.28]	0.02
R5–R20 kPa·L ⁻¹ ·s ⁻¹	0.05 [0.01–0.12]	0.03 [0.0–0.04]	NS
X5 kPa·L ⁻¹ ·s ⁻¹	–0.12 [–0.20––0.08]	–0.09 [–0.13––0.07]	NS
AX kPa·L ⁻¹	0.27 [0.21–1.15]	0.17 [0.13–0.38]	NS
Pulmonary function			
FVC % pred	102.6 [91.5–107.2]	111.0 [99.9–122.4]	NS
FEV1 % pred	94.4 [91.2–98.3]	106.9 [92.0–116.1]	NS
FEV1/FVC %	75.8 [71.0–86.1]	81.3 [75.4–90.2]	NS
FEV1/FVC <70%	1 [13]	2 [5]	NS
PEF % pred	80.4 [58.6–94.4]	101.6 [86.2–113.8]	0.01
MMF % pred	67.7 [45.5–91.8]	81.0 [56.5–104.9]	NS
FRC % pred	105.0 [97.5–116.8]	114.4 [95.1–121.4]	NS
RV % pred	117.3 [103.7–139.8]	101.8 [83.3–123.8]	NS
TLC % pred	101.2 [89.9–123.4]	101.9 [87.5–115.6]	NS
RV/TLC %	29.8 [24.9–35.9]	30.9 [25.3–33.8]	NS
D _{LCO} % pred	77.4 [64.4–86.3]	78.1 [68.6–89.6]	NS
eNO	29.4 [18.2–47.7]	21.5 [16.8–124.0]	NS
HRCT analyses			
Ai mm ²	22.3 [15.7–33.2], n=7	23.7 [18.0–34.4], n=36	NS
Ai/BSA ×10 ⁻⁶	13.6 [10.9–18.5], n=7	14.2 [11.6–19.9], n=36	NS
WA %	60.5 [55.3–66.5], n=5	56.5 [52.7–60.8], n=33	NS
Atr mm ²	225.2 [176.5–257.5], n=7	249.1 [214.0–319.4], n=37	NS
Atr/BSA ×10 ⁻⁶	114.3 [98.6–168.2], n=7	156.1 [133.6–189.9], n=37	NS
BAL	n=6	n=30	
Recovered cells ×10 ⁵ mL ⁻¹	1.80 [1.35–2.40]	1.90 [1.48–2.03]	NS
Lymphocytes %	39.9 [25.2–59.6]	25.4 [13.9–44.2]	NS
Eosinophils %	0.3 [0.0–1.1]	0.0 [0.0–0.0]	NS
CD4/CD8 ratio	2.90 [2.71–11.2]	5.90 [4.33–7.53]	NS

Data are presented as median [interquartile range] or n (%), unless otherwise stated. mMRC: modified Medical Research Council dyspnoea scale; ACE: angiotensin-converting enzyme; sIL-2R: soluble interleukin-2 receptor; KL-6: Krebs von den Lungen-6; IgE: immunoglobulin E; R5: resistance at 5 Hz; R20: resistance at 20 Hz; X5: reactance at 5 Hz; AX: area of reactance; FVC: forced vital capacity; FEV1: forced expiratory volume in 1 s; PEF: peak expiratory flow rate; MMF: maximum mid-expiratory flow rate; FRC: functional residual capacity; RV: residual volume; TLC: total lung capacity; D_{LCO}: diffusing capacity of the lung for carbon monoxide; eNO: exhaled nitric oxide; HRCT: high-resolution computed tomography; Ai: luminal area of right apical bronchus; BSA: body surface area; WA: wall area of right apical bronchus; Atr: tracheal luminal area; BAL: bronchoalveolar lavage; NS: not significant. [#]: theoretical range 1–5.

AHR status. Likewise, PFTs and F_{eNO} levels did not differ significantly between the two groups, except for the percentage of predicted peak expiratory flow rate. Neither HRCT indices nor BAL findings differed significantly by AHR status.

Of the 42 patients whose long-term outcomes (≥ 12 months) were available, seven were treated with systemic corticosteroids or an immunosuppressive agent for extrapulmonary lesions. Systemic corticosteroids were started for pulmonary sarcoidosis in another patient just after the comprehensive evaluation of airway. Of the remaining 34 patients in whom the natural course of pulmonary sarcoidosis was observed without systemic treatment, six (18%) experienced pulmonary disease progression after the median (interquartile range) observational period of 41.5 (12–78) months ($\geq 10\%$ decline in forced vital capacity compared to baseline, $\geq 10\%$ decline in forced expiratory volume in 1 s (FEV₁) during the period). Demographic data and serum biomarkers at diagnostic baseline were similar between patients with progressive and stable disease. Among IOS and PFT indices, median (interquartile range) X5 and AX were significantly lower and higher in patients with progressive disease, respectively (X5: -0.13 (-0.72 – -0.11) *versus* -0.09 (-0.11 – -0.06), $p=0.01$; AX: 0.40 (0.26 – 3.59) *versus* 0.16 (0.11 – 0.27); $p=0.01$), whereas PFT indices were comparable. Exhaled nitric oxide, HRCT indices and BAL findings did not differ between the groups.

We have shown that increased respiratory resistance is a better indicator for AHR in patients with early-stage pulmonary sarcoidosis than airflow limitation on PFTs. In a study of adult asthmatics, methacholine-induced dyspnoea and chest tightness did not correlate with changes in spirometry but did correlate with those in IOS [12]. Mildly asthmatic children have also displayed significantly higher IOS indices at baseline, relative to healthy controls, with no differences in FEV₁ [13]. Another study of AHR in sarcoidosis revealed that decreased FEV₁ predicted AHR although it did not include IOS measurements [14]. These results and our data both indicate the utility that IOS adds to even normal or mildly obstructive PFT profiles. Subjective respiratory symptoms such as cough, wheezing or dyspnoea on exertion were also typical accompaniments of AHR-positive sarcoidosis in our study. Given the low rate of airflow limitation in sarcoidosis [2], IOS may well serve as a screening tool for AHR in patients with early sarcoidosis, particularly if they have respiratory symptoms.

We have also demonstrated that decreased X5 and increased AX at the time of diagnosis are predictive for progression of pulmonary disease in patients with sarcoidosis. Although the determinants of X5 and AX values in IOS have not been fully defined, IOS may detect early airway disease and subtle parenchymal fibrosis without forced manoeuvres before these lesions have subsequently caused serious functional impairments on PFTs [4]. However, ventilation and lung volume may have affected these indices of IOS [10].

Some limitations of this study should be mentioned. First, the number of study subjects was small, and the follow-up period may not be enough to address the long-term outcome. Sarcoidosis was diagnosed only clinically without pathological confirmation in more than half of the subjects, which may have biased the study population. Another limitation was conducting this study at a single institute in Japan. Ethnic differences in clinical manifestations of sarcoidosis are well-known [15], and thus the prevalence of AHR and the long-term progression of pulmonary sarcoidosis may vary among differing ethnic groups. Finally, our patients did not receive uniform treatment, which may have skewed long-term outcomes of pulmonary lesions.

Despite these limitations, IOS has the potential to provide significant insight on AHR in early-stage pulmonary sarcoidosis. Reactance values may specifically predict the long-term progression of pulmonary involvement. Serial IOS measurements may also be useful to monitor the disease course.

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